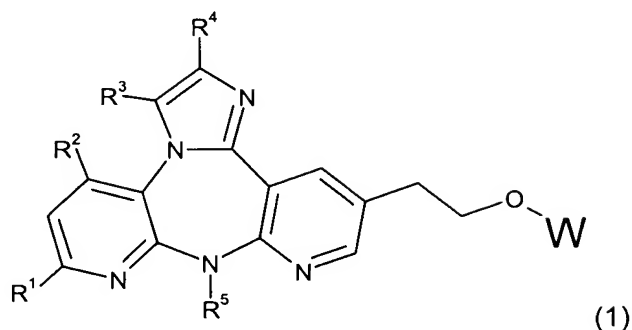


ABSTRACT

Compounds represented by formula 1:

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wherein **R**¹ is H, halogen, (C₁₋₄)alkyl, O(C₁₋₄)alkyl, and haloalkyl; **R**² is H or methyl; **R**³ is H or (C₁₋₄)alkyl; **R**⁴ is H or (C₁₋₄)alkyl; **R**⁵ is (C₁₋₄)alkyl, (C₁₋₄)alkyl(C₃₋₇)cyclo-alkyl or (C₃₋₇)cycloalkyl; and **W** is a fused phenyl-5 or 6-membered heterocycle having one or two heteroatoms selected from N or S; or **W** is phenyl, 1,1'-biphenyl, 2, 3-dihydro-1*H*-indene, 1, 2, 3, 4-tetrahydronaphthyl, or naphthyl; said **W** being optionally substituted with (C₁₋₄)alkyl, which in turn can be optionally substituted with a carboxy or (C₁₋₄)alkoxycarbonyl, or a salt or ester thereof. The compounds have inhibitory activity against Wild Type, single and double mutant strains of HIV.

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